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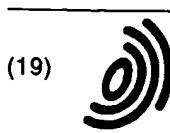
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(54) Process for making an acrylate coating

Verfahren zur Herstellung einer Acrylatbeschichtung

Procédé pour fabriquer un revêtement en acrylate

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(56) References cited:
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Description

[0001] This invention relates to deposition of barrier films for inhibiting penetration by oxygen or other gases. Such materials may be employed for fabricating high voltage capacitors. Techniques for acrylate deposition and adhesion are described.

[0002] Many products, including many food products, are packaged in thin plastic sheet bags or the like. The thin films are desirably resistant to permeation by oxygen, water vapor and odorous gases.

[0003] Such barrier films are commonly made of costly plastics because less costly films are too permeable to oxygen or water to give a long shelf life. Reduced cost barrier films are highly desirable.

[0004] There are many products that are packaged in plastic bottles, tubes or vials which also need protection from oxidation or contamination by environmental gases or which must be in impermeable containers so that components of the contents are not lost by diffusion through the containers.

[0005] US Patent No. 4 842 893 discloses a high-speed process for coating substrates with thin organic coatings in a vacuum. The process involves cooling the support (drum) to a temperature in the range of 20°C to 80°C and condensing an evaporated acrylate monomer of a molecular weight of 150-1000 on a substrate secured to the cooled drum.

[0006] Electrical capacitors are used for storing energy in a variety of applications. Operating voltages for such devices range from a few volts, such as those used in miniature electronic circuits, to thousands of volts, such as those used in power utility applications.

[0007] For low and medium voltages (generally below 600 volts) metallized sheet dielectrics are commonly used. High voltage capacitors are generally constructed of aluminium foil electrodes and sheet dielectric material such as a thermoplastic polypropylene and polyester film. Capacitors intended for service at more than about 600 volts are completely impregnated with a dielectric liquid with good gas absorbing properties. Typical dielectric liquids with good gas absorbing properties are PXE (phenyl xylyl ethane), MIPB (mono isopropyl biphenyl), DOP (diethyl phthalate), castor oil, polypropylene glycol and mineral oil.

[0008] High voltage capacitors have not been able to take advantage of the self healing properties of metallized electrodes. The dielectric liquid causes swelling of the thermoplastic sheets, which causes cracking of the thin film metal electrode and an open circuit quickly develops. It would be desirable to have wound capacitors made with metallized film for higher voltages where self healing can occur, and often where it is necessary that the capacitor be liquid filled.

[0009] There is, therefore, a method for forming a substrate coated with an acrylate layer defined by the features of claim 1. The substrate is first chilled to a temperature between 0°C and 15°C outside of a deposition

station. The chilled substrate is passed through a deposition station using a coating drum cooled to a temperature between 0°C and 15°C. At the deposition station an acrylate monomer having an average molecular weight in the range of from 200 to 600 is evaporated and condensed on the substrate as an acrylate film. The acrylate monomer film is polymerized for forming a polymerized acrylate layer.

[0010] There is further a method which is defined by the features of claim 7.

[0011] FIG. 1 illustrates schematically a metallized sheet capacitor.

[0012] FIG. 2 illustrates in three transverse cross sections materials suitable for winding a capacitor, FIG. 2a illustrating two coated sheets as used in a capacitor, FIG. 2b illustrating a sheet coated on only one face, and FIG. 2c illustrating a sheet with the coating sequence reversed.

[0013] FIGS. 3, 4 and 5 illustrate, in fragmentary transverse cross section, other embodiments of material suitable for winding capacitors.

[0014] FIG. 6 is a schematic illustration of coating apparatus for forming such a capacitor sheet material.

[0015] FIG. 7 is a graph with the axes representing acrylate shrinkage and layer thickness, respectively.

[0016] FIG. 8 is a graph illustrating condensation efficiency as a function of temperature.

[0017] FIG. 9 is a fragmentary cross section of a capacitor made with metallized sheet coated with a polymerized acrylate.

[0018] FIG. 10 illustrates in transverse cross section a coated polypropylene with low oxygen permeability.

[0019] FIG. 11 illustrates in transverse cross section a coated thermoplastic substrate with low oxygen permeability.

[0020] FIG. 12 illustrates an exemplary container constructed according to principles of this invention.

[0021] FIG. 13 is a schematic illustration of apparatus for coating substrates by spraying.

[0022] FIG. 14 is a schematic illustration of coating apparatus for forming a barrier sheet material.

[0023] An exemplary capacitor 10 comprises a can 11, the end of which is cut away in FIG. 1 to show the roll of capacitor material 12 inside the can. The layers of material forming the capacitor are wound into a cylindrical roll. In effect, the roll of capacitor material comprises alternating layers of metal and dielectric coiled in a tight spiral. The metal layers form the electrodes of the capacitor and the dielectric material between the electrodes permits accumulation of charge in the capacitor. Such a capacitor, intended for use at higher voltages, may be filled with a dielectric liquid such as those mentioned above.

[0024] FIG. 2 illustrates in three fragmentary transverse cross sections exemplary materials for winding such a capacitor. A sheet suitable for winding a capacitor has a thermoplastic substrate 13 such as thin polypropylene or polyester sheet which has a stable dielec-

tric constant and high dielectric strength. A layer of metal 14 which may be vacuum deposited or sputtered is formed on at least one face of the substrate. A layer of polymerized acrylate 16 is deposited either over or under the metal layer as described hereinafter. In the embodiment illustrated in FIG. 2a, another layer of polymerized acrylate 17 is formed on the opposite face of the substrate.

[0025] The two sheets of material suitable for forming a capacitor are illustrated in FIG. 2a as they are sometimes used in capacitors. Each sheet has a portion of the metallized layer 14 exposed along an edge of the sheet. That is, a portion of the metallized layer is not covered by an insulating layer of acrylate in this embodiment. These exposed edges of the metal layer are at opposite edges of adjacent sheets in the capacitor and provide an area to which electrical contact can be made so that alternate sheets of the capacitor material form the opposite plates of the capacitor.

[0026] FIG. 2b illustrates another embodiment of sheet for forming a rolled capacitor. Here a layer of metal 14 is deposited on the substrate 13 and a layer of polymerized acrylate 16 is formed over the metal layer. The entire metal layer is covered with acrylate. Only one face of the substrate has a coating of acrylate.

[0027] The embodiment illustrated in FIG. 2c reverses the layers on the substrate. In this embodiment there is a layer of polymerized acrylate 16 formed directly on the substrate 13. In direct contact with and over the acrylate layer is a deposited layer of metal 14. Again, there is no acrylate layer on the opposite face of the substrate. The metal may extend all the way to each edge of the sheet or may be spaced a short distance from the edge of the sheet along one edge.

[0028] FIG. 3 illustrates a variation of the capacitor material having a layer of polymerized acrylate 18 on each face of a thermoplastic substrate 19. A layer of metal 21 is deposited on top of one of the acrylate layers.

[0029] FIG. 4 illustrates another slight variation from an embodiment as illustrated in FIG. 3, having a layer of polymerized acrylate 22 on each face of a substrate 23. A layer of metal 24 is deposited over at least one of the acrylate layers. A third layer of polymerized acrylate 26 is formed over the layer of metal. Any of these variations may be used for forming capacitors.

[0030] It is not always required that a capacitor be formed with both electrodes being metallized sheets of dielectric material. Thus, for example, as illustrated in FIG. 5, a capacitor may have one electrode formed on a thermoplastic substrate 27. A layer of polymerized acrylate 28 is formed on each face of the substrate. A layer of metal 29 is deposited for forming the one electrode. The other electrode in such a capacitor comprises a sheet of aluminum foil 31. That electrode may be separated from an adjacent layer of metal (not shown) on another substrate via a sheet 32 of dielectric materials such as polypropylene.

[0031] As mentioned above, the polypropylene and

polyester materials are desirable as dielectrics for capacitors since they have a stable dielectric constant and high dielectric strength. If a metallized polypropylene, for example, is tried in a liquid filled capacitor, the thermoplastic apparently swells and ruptures the thin metallized layer. By having a very thin layer of polymerized acrylate in direct contact with the metallized electrode, the damage caused by dielectric liquid can be avoided.

[0032] It is believed that the thermoset acrylate supports or stabilizes the metal film and prevents cracking despite swelling of the film substrate.

[0033] An exemplary barrier material as illustrated in FIG. 11 comprises a thermoplastic substrate 110 having a cross-linked acrylate layer 111 on one face. The acrylate layer is overlain by a layer 112 of oxygen barrier material. A final cross-linked acrylate layer 113 is optionally formed over the oxygen barrier layer.

[0034] It has been discovered that polyethylene, polypropylene, polyester or nylon substrates with thin surface coatings of crosslinked acrylate have very low oxygen and water permeability when combined with a metal or inorganic barrier coating. Typical sheets for packaging foodstuffs include metallized polypropylene, nylon or polyester sheet. Metallized nylon has an oxygen permeability of about 0.05 ml/100 in²/hour (ml/645 cm²/hour) as measured with a Mocon Oxtran System available from Modern Controls, Minneapolis, Minnesota.

[0035] Metallized polyester has a typical oxygen permeability of about 0.08. Metallized polypropylene, on the other hand, has an oxygen permeability of about 2.5 and is not usually suitable for packaging where low oxygen permeability is important.

[0036] It is believed that the high oxygen permeability of metallized polypropylene is due to the inherent surface roughness of the polypropylene sheet. Nylon and polyester sheets are considerably smoother and have a higher temperature capability than polypropylene. A metal coating of uniform thickness can be readily applied as a good oxygen barrier. A layer of acrylate about 1/2 to one micrometer thick is adequate for smoothing the surface for producing a low oxygen permeability.

[0037] Sheet polypropylene without any coating may have an oxygen permeability of about 100. However, if a layer of aluminum is applied to a surface of a polypropylene sheet substrate, the oxygen permeability decreases to about 2.5. Surprisingly, when an acrylate layer only about one micrometer thick is formed on the polypropylene and then covered with a layer of metal, the oxygen permeability drops to about 0.05, a value lower than metallized polyester. It is hypothesized that the film of liquid acrylate monomer deposited on the surface of

the polypropylene has a smooth, high temperature surface and the surface remains smooth when the acrylate is polymerized. The metallized layer can then form a good oxygen barrier.

[0037] First, a layer of acrylate monomer is deposited on the substrate and crosslinked. The acrylate layer is then coated with an oxygen barrier layer of SiO_x or Al_2O_3 , both of which have good resistance to oxygen permeability. The high temperature resistance of the crosslinked acrylate layer permits the notably higher temperature deposition of silicon oxide or aluminum oxide on the thermoplastic substrate. Other deposited barrier films may be formed of nitrides or silicides that can be vacuum or vapor deposited. For example, silicon nitride, aluminum nitride, or titanium nitride may be used. The materials need not be stoichiometric compounds and often are stoichiometric with excess metal.

[0038] A still greater surprise occurs when another polymerized acrylate layer is formed over the oxygen barrier layer. The permeability through a polypropylene barrier material drops to about 0.002 which is appreciably better than the oxygen permeability of metallized nylon. It is hypothesized that the second acrylate layer may protect the metallized layer and assure retention of the oxygen barrier properties of the metal.

[0039] A preferred composite material with low oxygen permeability has a layer of polymerized acrylate, a layer of barrier material such as SiO_x , Al_2O_3 , silicon nitride, titanium nitride or aluminum nitride and another layer of polymerized acrylate on a thermoplastic substrate. The layers of acrylate reduce permeability dramatically and the layer overlying the barrier material protects the barrier material from mechanical damage and also provides a surface suitable for printing.

[0040] The substantial improvement in oxygen permeability is believed to be attributable to formation of a liquid film of monomer on the surface of the polypropylene, followed by cross linking of the polyfunctional acrylate. Adding a second layer of acrylate monomer which is polymerized in situ is believed to rectify any defects in the underlying layers and provide an additional thickness of material with inherently low oxygen permeability. Redundant layers minimize pinhole leakage.

[0041] A particularly useful barrier sheet may be formed with multiple layers of acrylate and oxygen barrier. Several thin layers of such materials are more effective than a couple thicker layers with the same total thickness. For example, a sheet may be coated with a layer of cross linked acrylate about 0.5 micrometers thick followed by a layer of metal or other oxygen barrier about 0.1 micrometer thick. Subsequent layers of acrylate and metal are built up, preferably with the final layer being acrylate. The thin layers of acrylate between layers of metal reduce the lateral flow of oxygen in the layers and minimize permeability. Such an arrangement is particularly useful as an oxygen barrier on bottles or vials.

[0042] Although it is generally preferred to deposit a

layer of acrylate on the substrate, followed by a layer of oxygen barrier material, oxygen permeability can be sharply reduced with a layer of metal deposited directly on the substrate, followed by a layer of acrylate.

5 [0043] The polymerized acrylate layer is believed to be beneficial for a number of other reasons. As a thermoset material, it has higher temperature resistance than the thermoplastic substrate. In the coating process, the product is subjected to elevated temperature processing such as metallizing, plasma treatment and the like. A cured acrylate coating would not have vapor emissions and may seal the surface and inhibit emission of materials from a thermoplastic substrate.

10 [0044] It is particularly preferred to vaporize an acrylate prepolymer and deposit it on a substrate in a vacuum system since this technique serves to refine the acrylate. In effect, the vaporization and deposition process is a vacuum distillation which removes volatiles to the vacuum pumps and deposits only higher molecular weight acrylates on the substrate. Removal of volatiles is desirable for subsequent high temperature processing of the material such as in deposition of metal or other inorganic barrier films.

15 [0045] After any of these deposition techniques, the monomer film is irradiated with ultraviolet or an electron beam to cause polymerization of the acrylate to form a monolithic crosslinked layer. Polymerization by irradiation is a conventional practice and the electron flux required or wavelength and total flux of ultraviolet used are commonly known. A photoinitiator may be included in the acrylate for facilitating polymerization by ultraviolet radiation.

20 [0046] A suitable apparatus for coating the substrate with acrylate and metal layers is illustrated schematically in FIG. 6. All of the coating equipment is positioned in a conventional vacuum chamber 36. A roll of polypropylene, polyester or nylon sheet is mounted on a payout reel 37. The sheet 38 forming the substrate is wrapped around a first rotatable drum 39, around a second rotatable drum 40, and fed to a take-up reel 41. Idler rolls 42 are employed, as appropriate, for guiding the sheet material from the payout reel to the drums and to the take-up reel.

25 [0047] A flash evaporator 43 is mounted in proximity to the drum at a first coating station. The flash evaporator deposits a layer or film of acrylate monomer on the substrate sheet as it travels around the drum. After being coated with acrylate monomer the substrate sheet passes an irradiation station where the acrylate is irradiated by a source 44 such as an electron gun or source of ultraviolet radiation. The radiation or electron bombardment of the film induces polymerization of the acrylate monomer.

30 [0048] The sheet then passes a metallization station 46 where a coating of metal for an electrode is applied by vacuum metallizing or sputtering. The sheet then passes another flash evaporator 47 where another layer of acrylate monomer is deposited for forming a protec-

tive layer over the metal. This layer of monomer is cured by irradiation from an ultraviolet or electron beam source 48 adjacent the drum. Depending on whether a layer of acrylate is above or below the metal layer, either of the evaporators 43 or 47 may be used. Clearly, if the metal layer is to be sandwiched between layers of acrylate, both evaporators and their respective radiation sources are used.

[0049] An exemplary process for coating a container can be described as follows. The container is injection molded or blow molded in a conventional manner from a conventional thermoplastic material. Preferably the container is then flame treated for activating and smoothing the surface. It has been found that adhesion of an acrylate layer on the substrate is enhanced by activating the surface by plasma or flame treating. High temperature air may also be used.

[0050] In a typical production line, a row of containers 114 are moved successively through a flame treating station, a coating station and a curing station. In the flame treating station, the containers are bathed in the flames from a plurality of propane or natural gas torches 116. In addition to activating the surface for enhanced adhesion, the flame treating can significantly smooth the surface of the container to assure that there is complete coverage by subsequent coatings. The flame is applied to the surface with sufficient intensity to soften or melt a thin surface layer on the container.

[0051] The flame treatment melts and rounds off any molding flash and smooths mold marks on the container so that the coating can bridge surface irregularities. "Melting" may almost be considered a misnomer since the thermoplastic materials are effectively supercooled liquids. Thus, melting is considered to be sufficient softening of the surface for smoothing irregularities.

[0052] Such an acrylate layer is applied in a coating station where one or more nozzles 17 sprays a thin coating of acrylate monomer onto the surface. Such a sprayed coating may be in the order of from one to twenty micrometers thick, for example. The acrylate sprayed onto the surface may be a low viscosity monomer or if desired, a monomer and/or a low molecular weight polymer may be combined with a solvent for spraying. In the event the acrylate is to be cross-linked by ultraviolet irradiation, a photoinitiator may be included in the sprayed composition.

[0053] An alternative to spraying the surface of the container with acrylate comprises dipping the container into a liquid acrylate composition.

[0054] Following the coating station, the containers pass a curing station where a plurality of ultraviolet lamps 18 irradiate the acrylate layer and cause cross-linking.

[0055] Following the application of an acrylate layer to the containers, an oxygen barrier layer is applied. Preferably this is by deposition in a vacuum chamber. A metal barrier layer, e.g. aluminum, may be applied by vacuum metallizing or sputtering. A layer of silicon ox-

ide, aluminum oxide, silicon nitride or other such material may be deposited by a plasma assisted chemical vapor deposition technique. For example, SiO_x may be deposited by a plasma vapor deposition process using an oxidizing or inert carrier gas. Aluminum oxide can be deposited by electron beam evaporation or preferably by evaporation of aluminum which is converted to an oxide in an oxygen plasma.

[0056] An alternative technique for depositing an acrylate layer is in a vacuum chamber. Suitable apparatus for coating a sheet substrate with acrylate and oxygen barrier layers is illustrated schematically in FIG. 14. All of the coating equipment is positioned in a conventional vacuum chamber 121. A roll of polypropylene, polyester or nylon sheet is mounted on a pay-out reel 122. The sheet 123 forming the substrate is wrapped around a rotatable drum 124 and fed to a take-up reel 126. Idler rolls 127 are employed, as appropriate, for guiding the sheet material from the payout reel to the drum and to the take-up reel.

[0057] A flash evaporator 128 is mounted in proximity to the drum at a first coating station. The flash evaporator deposits a layer or film of acrylate monomer on the substrate sheet as it travels around the drum. After being coated with acrylate monomer the substrate sheet passes an irradiation station where the acrylate is irradiated by a source 129 such as an electron gun or source of ultraviolet radiation. The UV radiation or electron bombardment of the film induces polymerization of the acrylate monomer.

[0058] The sheet then passes a deposition station 131 where a coating of oxygen barrier material is applied by plasma deposition, vacuum deposition or the like. The sheet then passes another flash evaporator 132 where another layer of acrylate monomer is deposited for forming a protective layer over the oxygen barrier. This layer of monomer is cured by irradiation from an ultraviolet or electron beam source 133 adjacent the drum. The coated sheet is then wrapped up on the take-up reel 126.

[0059] Evaporation of the monomer is preferably from flash evaporation apparatus 129, 132 as described in U.S. Patents Nos. 4,722,515, 4,696,719, 4,842,893, 4,954,371 and/or 5,097,800. These patents also describe polymerization of acrylate by radiation. In such flash evaporation apparatus, liquid acrylate monomer is injected into a heated chamber as droplets. The elevated temperature of the chamber vaporizes the droplets to produce a monomer vapor. The monomer vapor fills a chamber with a longitudinal slot forming a nozzle through which the monomer vapor flows.

[0060] Two styles of evaporator are suitable. In one of them, the orifice for injecting droplets and flash evaporator are connected to one end of the nozzle cylinder. In the other style, the injector and flash evaporator section is attached in the center of the nozzle chamber like a T.

[0061] It is often found desirable to plasma treat the

surface to be coated immediately before coating. A conventional plasma gun 134 is positioned in the vacuum chamber upstream from each of the flash evaporators 128 and 132 for activating the surface of the sheet on a continuous basis before monomer deposition. Conventional plasma generators are used.

[0062] Surface treatment within the vacuum chamber is of substantial importance for all surfaces, not just the surface of the raw film. Referring again to FIG. 6, a conventional plasma gun 52 is positioned in the vacuum chamber upstream from each of the flash evaporators 43 and 49 for activating the surface of the sheet on a continuous basis before monomer deposition. Another plasma gun 52 is provided immediately before the vacuum metallizing station 46. Conventional plasma generators are used.

[0063] In an exemplary embodiment the plasma generator is operated at a voltage of about 500 to 1000 volts with a frequency of about 50 KHz. Power levels are in the order of 500 to 3000 watts. For an exemplary 50 cm wide film traveling at a rate of 30 to 90 meters per minute, around 500 watts appears appropriate.

[0064] An analogous system may be employed for coating containers with layers of acrylate and barrier material. The containers are moved successively through an acrylate evaporation and condensation station and a deposition station. If two layers of acrylate are used the second acrylate may be applied by way of the same flash evaporator or by way of a second similar flash evaporator.

[0065] The acrylates used for forming the cross-linked coatings on the thermoplastic substrate differ somewhat depending on the technique used for depositing the coating. The acrylates used for dipping or spraying are similar and it is not necessary that the acrylate is a monomer. Generally, the acrylates used are blends of high and low molecular weight materials to yield the desired viscosity of the composition for dipping or spraying. Monomers with molecular weights in the order of 150 up to partially polymerized materials have a molecular weights in the order of 20,000 may be blended to obtain a low viscosity blend. There should be polyfunctional acrylates in the blend so that there is extensive cross-linking.

[0066] In the event the acrylate layers are applied by the evaporation and condensation technique, the range of suitable acrylates is more restricted. These acrylate resins are generally monomers having a molecular weight in the range of from 150 to 600. Preferably, the monomers have a molecular weight in the range of from 200 to 400. Higher molecular weight fluorinated acrylates or methacrylates may be equivalent to these lower molecular weight materials and also be used for forming a deposited acrylate layer. For example, a fluorinated acrylate with a molecular weight of about 2000 evaporates and condenses similar to a non-fluorinated acrylate having a molecular weight in the order of 300. The acceptable range of molecular weights for fluorinated

acrylates is about 400 to 3000. Fluorinated acrylates include monoacrylates, diacrylates, and methacrylates. Chlorinated acrylates may also be useful.

[0067] If the molecular weight is below about 150, the monomer is too volatile and does not condense well for forming a monomer film. If the molecular weight is more than about 600 the monomer does not evaporate readily in the flash evaporator at temperatures safely below the decomposition temperature of the monomer.

[0068] When the monomers polymerize, there may be shrinkage of the film. Excessive shrinkage may cause poor adhesion of the layer on the substrate. Shrinkage up to about 15 to 20% can be tolerated in the thin acrylate layers. However, it is preferred that the shrinkage be less than 10% for reliable coating adhesion.

[0069] A typical monomer used for flash evaporation includes an appreciable amount of diacrylate and/or triacrylate to promote crosslinking. Blends of acrylates may be employed for obtaining desired evaporation and condensation characteristics and adhesion, and for controlled shrinkage of the deposited film during polymerization.

[0070] Suitable monomers are those that can be flash evaporated in a vacuum chamber at a temperature below the thermal decomposition temperature of the monomer and below a temperature at which polymerization occurs in less than a few seconds at the evaporation temperature. The monomers selected should also be readily capable of crosslinking when exposed to ultraviolet or electron beam radiation.

[0071] The monomer composition may comprise a mixture of monoacrylates and diacrylates. Triacrylates tend to be reactive and may polymerize at the evaporation temperatures. Generally speaking, shrinkage is reduced with higher molecular weight materials.

[0072] Generally, it is desirable that at least a major portion of the acrylate monomer evaporated is a poly-functional acrylate for crosslinking. Preferably, the acrylate comprises at least 70 percent polyfunctional acrylates such as diacrylate or triacrylate. If the degree of crosslinking is too low, the polymerized acrylate layer may not have adequate cure speed.

[0073] There are situations when less than half of the acrylate monomer comprises polyfunctional acrylates. For example, in a dry capacitor film one composition comprises about 80% monoacrylate and 20% diacrylate. The high proportion of monoacrylate is used since it has a high dielectric constant.

[0074] Preferably, the molecular weight of the acrylate monomer is in the range of from 200 to 400. If the molecular weight is less than about 200, the monomer evaporates readily, but may not condense quantitatively on the substrate without chilling of the substrate. If the molecular weight is more than about 400, the monomers become increasingly difficult to evaporate and higher evaporation temperatures are required.

[0075] Preferably, the acrylate monomer has a vapor pressure at 25°C in the range of from 1 to 20 microme-

ters of mercury. If the vapor pressure is less than about one micrometer, exceptionally high temperatures may be required to evaporate sufficient material for forming a coating on the substrate in reasonable coating time. High temperatures may lead to thermal decomposition or premature curing of the monomers. If the vapor pressure is higher than about twenty micrometers of mercury, condensation of the monomer to form a film on the substrate may have too low an efficiency for practical coating operations. Adequate efficiency may not be obtained until the surface of the substrate is cooled below the freezing point of the monomer, in which case the material may not polymerize properly.

[0076] There are at least five monoacrylates, ten diacrylates, ten to fifteen triacrylates and two or three tetraacrylates which may be included in the evaporated composition. Most preferably the acrylate comprises hexane diol diacrylate (HDDA) with a molecular weight of 226 and/or tripropylene glycol diacrylate (TRPGDA) with a molecular weight of about 300. Other acrylates may be used, sometimes in combination, such as monoacrylates 2-phenoxy ethyl acrylate (M.W. 192), isobornyl acrylate (M.W. 208) and lauryl acrylate (M.W. 240), epoxy acrylate RDX80095 made by Radcure of Atlanta, Georgia; diacrylates diethylene glycol diacrylate (M.W. 214), neopentyl glycol diacrylate (M.W. 212), propoxylated neopentyl glycol diacrylate (M.W. 328) and polyethylene glycol diacrylate, tetraethylene glycol diacrylate (M.W. 302), and bisphenol A epoxy diacrylate; and triacrylates trimethylol propane triacrylate (M.W. 296), ethoxylated trimethylol propane triacrylate (M.W. 428), propylated trimethylol propane triacrylate (M.W. 470) and pentaerythritol triacrylate (M.W. 298). Monomethacrylates isobornyl methacrylate (M.W. 222) and 2-phenoxyethyl methacrylate (M.W. 206) and dimethacrylates triethylene glycol dimethacrylate (M.W. 286) and 1,6-hexanediol dimethacrylate (M.W. 254) may also be useful, but may cure too slowly to be useful for high speed coating operations.

[0077] It has generally been considered that it is not feasible to evaporate high molecular weight acrylates because of their very low vapor pressure and high viscosity. Evaporated acrylate coatings have been restricted to low molecular weight monomers, generally below a molecular weight of about 400 and with low viscosity. Generally the viscosities are below 50 centistoke (1 Stoke = $1 \text{ cm}^2 \text{s}^{-1}$).

[0078] It has been found, however, that by mixing a very low and a very high viscosity material, flash evaporation, condensation and curing can be obtained. For example, a mixture of 70 percent tripropylene glycol diacrylate (TRPGDA) and 30 percent of beta carboxy ethyl acrylate (BCEA) has a viscosity of about 15 centistokes and can be readily evaporated, condensed and cured. The low viscosity component lowers the viscosity of the blend, which improves atomization in the evaporator and assists in the flash evaporation of the high viscosity acrylate.

[0079] When blends of high and low molecular weight acrylates are used, it is preferred that the weighted average molecular weight of the blend be in the range of from 200 to 600 and preferably up to about 400. This assures that there is good vaporization of the blend at reasonable temperatures in the evaporator.

[0080] Some examples of low molecular weight acrylates are hexane diol diacrylate, diethylene glycol diacrylate, propane diacrylate, butane diol diacrylate, tripropylene glycol diacrylate, neopentyl glycol diacrylate, phenoxyethyl acrylate, isobornyl acrylate and lauryl acrylate. Some examples of high molecular weight acrylates are bisphenol A diacrylate, BCEA, Radcure 7100 (an amine acrylate available from Radcure, Atlanta Georgia), Radcure 169, Radcure 170, acrylated and methacrylated phosphoric acid, Henkel 4770 (an amine acrylate available from Henkel Corporation, Ambler, Pennsylvania) and glycerol propoxy triacrylate.

[0081] Particularly preferred high molecular weight materials include BCEA which is acid in character and has a shrinkage of only about 4 percent upon curing. Another suitable material is an acrylate or methacrylate of phosphoric acid. One can also use acrylic acid in the composition, along with dimers, trimers and tetrameres of acidic acrylates or methacrylates. The molecular weight range of the acrylate may be extended by pre-heating the prepolymer before it is atomized into the vaporization chamber.

[0082] It has been found that the temperature of the substrate on which the monomer film is deposited can have a large influence on the efficiency of condensation. It is desirable to refrigerate the substrate before it is placed in the vacuum chamber. Good condensation efficiency can be obtained with monomers having a molecular weight of at least 200 with the substrate cooled to temperatures in the range from 0 to 15°C. A rack of containers to be coated can be removed from a refrigerator and placed in a vacuum chamber, pumped down and coated before the containers warm to unreasonably high temperatures.

[0083] The rotating drums are also cooled so that the substrate remains at a low temperature. When the sheet being coated is smooth and thin (generally less than 12 micrometers) good condensation efficiency can be obtained with monomers having a molecular weight of at least 200 with the coating drum cooled to temperatures in the range from 0 to 15°C.

[0084] Fig. 8 includes a series of data points showing measured condensation efficiencies of hexane diol diacrylate monomer as a function of the substrate temperature. HDDA is a relatively volatile monomer with a molecular weight about 212. A less volatile monomer such as tripropylene glycol diacrylate with a molecular weight of about 300 has higher condensation efficiency.

[0085] An extreme amount of cooling of the substrate must also be avoided. For example, HDDA freezes on the substrate when the coating drum temperature is below about 0°C. The frozen monomer cannot be polym-

erized. Thus, the cooling temperature must hold the surface of the substrate above the freezing point of the monomer but below the temperature at which the condensation efficiency decreases significantly.

[0086] In addition to precooling the roll of substrate the sheet can be cooled on the front surface preceding the evaporator. For example, the idler roll 42 between the payout reel and the first drum may be chilled for cooling the front surface of the substrate sheet before it reaches the first drum. Cooling the front face of the sheet, i.e. the face on which deposition is to occur can permit faster processing and permits cooling of sheets with a rough reverse surface.

[0087] Precooling of a sheet of material on which the acrylate is to be deposited can be significant for applications in addition to preparation of sheet material for winding capacitors. For example, there are times that it is desirable to coat paper with an acrylate and the rough surface of paper does not lend itself to cooling from the back surface on a rotating drum. In such a case, the flexible paper may be passed over a chilled roll to cool the front face of the paper. An acrylate is then deposited on the same face of the paper as was just cooled before the face heats any significant amount.

[0088] There may be embodiments where it is sufficient to deposit an oxygen barrier layer directly on the substrate and apply an acrylate layer over the oxygen barrier material. For example, when the thermoplastic substrate has been flame treated to smooth the surface sufficiently that a thin oxygen barrier material can bridge over any surface irregularities, the oxygen barrier material may be deposited directly on the flame treated substrate. An acrylate layer may then be applied over the oxygen barrier material to protect the barrier and further reduce permeability.

[0089] It is found particularly desirable to provide a protective crosslinked acrylate coating over a deposited layer of metal such as aluminum. If an aluminum layer is applied to a sheet substrate which is rolled for later use or which is passed over a roller contacting the surface, the aluminum may be abraded off of higher asperities on the surface. A sheet coated with aluminum and protected with an overlying crosslinked acrylate coating having a thickness of as little as 0.1 micrometer, the pin-hole density can be maintained as low as 10 pinholes per cm².

[0090] When films used in packaging food products, cigarettes or many other items, the presence of an acrylate odor is unacceptable. Any such odor can be eliminated by curing any residual acrylate monomer on the film before it is removed from the vacuum chamber.

[0091] An electron gun 53 is mounted in the vacuum chamber between the final idler roll 42 and the take-up reel 41. Electron bombardment from the gun polymerizes any residual acrylate monomer on the surfaces of the film before it is rolled up. By irradiating in the converging space between the film on the take-up reel and the film advancing from the idler roll both faces of the film can

be irradiated with a single electron gun.

[0092] A sheet capacitor is often constructed by winding two sheets of metallized plastic sheet together, with the metallized layer on one sheet being one electrode of the capacitor and the metallized layer on the other sheet being the other electrode. The two sheets are offset from each other so that at each end of the roll the edge of one sheet extends beyond the edge of the other sheet. The electrical contact to the two electrodes is made by way of the protruding edge of each electrode at opposite ends of the roll.

[0093] Molten zinc or lead-tin solder is then sprayed onto the ends of the roll (a process called Schooping) to establish electrical contact with the metallized layer.

15 A connecting lead is either welded or soldered to the relatively thick contact material. The zinc or solder Schooping apparently enters between the metallized plastic sheets and may melt a portion of the sheet to make mechanical and electrical contact to the thin metallized layer on the sheet.

[0094] It is preferred to have a layer of polymerized acrylate overlying the metallized layer of the capacitor material. A layer of acrylate in the range of from about 0.1 to 1 micrometer thick has been found sufficient to prevent corrosion of a zinc electrode in a high humidity environment. Furthermore, the overlying layer of acrylate improves the scratch resistance of the metallized electrode. The electrode may be abraded during slitting of the metallized sheet or winding a capacitor. An overlying acrylate layer effectively prevents damage from such scratching. The overlying layer of acrylate also helps confine the tiny arc that occurs adjacent to a short, which increases its temperature for more efficiently evaporating the metal from around the shorted area.

20 30 35 This, and the presence of oxygen in the acrylate material, enhance the self-healing of the capacitor.

[0095] FIG. 9 illustrates in a fragmentary transverse cross section a few layers of a preferred capacitor. Each of the layers of the capacitor comprises a substrate sheet 56 having a metallized layer 57 on one face. A layer 58 of polymerized acrylate is deposited over each metallized layer. A second acrylate layer 59 may be deposited on the opposite face of each dielectric sheet 56 for isolating the sheet from dielectric capacitor liquid. A metal contact electrode 60 is sprayed onto each end of the roll of metallized sheets.

[0096] The edges of the two sheets forming the capacitor are offset from each other. Thus, at one end of the capacitor, the edge of one sheet A extends beyond the edge of the intervening sheet B which is recessed. At the opposite end of the capacitor the recessed sheet B extends beyond the edge of the interleaved sheet A. This permits electrical contact to be made to the metallized layers on the respective sheets at opposite ends 40 45 50 55 of the capacitor.

[0097] In this embodiment the metallized layer on each sheet extends to one edge of the sheet and does not extend all the way to the opposite edge of the sheet.

Thus, for example, as illustrated at the left edge of FIG. 9, the metallized layer extends all the way to the edge of the protruding sheets A. On the other hand, the metallized layer on the recessed sheets B stop short of the edge of the sheet. Because of this, when the contact metal 60 is sprayed onto the end of the roll, electrical contact can be made to the metallized layer on the protruding sheet and the metallized layer on the recessed sheet is electrically insulated by the plastic material.

[0098] Surprisingly, despite the substantially complete enclosure of the metallized layer by acrylate and polypropylene plastic, good electrical contact is made when the electrical contact is sprayed onto the end of the roll. It is believed that a small amount of thermoplastic dielectric sheet melts, thereby assuring good electrical contact with the metallized layer. Polypropylene has a melting temperature of about 165°C, and the cross linked acrylate does not melt. The metallized layer is, in effect, supported by the polymerized acrylate which resists higher temperatures.

Claims

1. A method for forming a substrate (123) coated with an acrylate layer comprising the steps of evaporating an acrylate monomer, having an average molecular weight in the range of from 200 to 600, at a deposition station (131), condensing the acrylate monomer on the substrate (123) as an acrylate film, polymerizing the acrylate film for forming a polymerized acrylate layer **characterised in that** the substrate (123) is chilled to a temperature between 0°C and 15°C outside of the deposition station (131) and the chilled substrate is passed through the deposition station (131) using a coating drum cooled to a temperature between 0°C and 15°C. 25
2. A method according to Claim 1, wherein the substrate comprises a flexible sheet and comprising the steps of:

engaging a face of the flexible sheet with a chilled roller (42); and depositing the acrylate on the same face that engaged the chilled roller (42). 30
3. A method according to either one of Claims 1 to 2, wherein the substrate comprises a plastic film and comprising the steps of:

chilling a roll of plastic film,
unrolling the plastic film,
engaging the plastic film with a chilled drum, and
rotating the drum past the deposition station (131). 35
4. A method according to either one of Claims 1 or 2, 5 wherein the substrate comprises a flexible sheet film and comprising the steps of:

engaging a first face of the sheet with a chilled roller (42);
thereafter engaging the other face of the sheet with a chilled drum; and
rotating the drum past the deposition station for depositing acrylate on the first face. 40
5. A method according to any one of Claims 1 to 4, wherein the acrylate monomer is a polyfunctional acrylate prepolymer and the evaporating step comprises the steps of:

preheating the polyfunctional acrylate prepolymer to a temperature above ambient temperature and lower than a temperature where the prepolymer polymerizes;
atomizing the preheated prepolymer into a heated vaporization chamber in a vacuum system; and
vaporizing the prepolymer in the vaporization chamber and discharging the vaporized prepolymer into the vacuum system. 45
6. A method according to any one of Claims 1 to 4, wherein the acrylate is a blend of acrylate monomers comprising at least one monomer having a molecular weight in the range of from 200 to 400 and an acrylate having a molecular weight of more than 600. 50
7. A method for forming a substrate (123) coated with an acrylate layer comprising the steps of evaporating an acrylate monomer, having an average molecular weight in the range of from 400 to 3000, at a deposition station (131), condensing the acrylate monomer on the substrate (123) as an acrylate film, polymerizing the acrylate film for forming a polymerized acrylate layer **characterised in that** the substrate (123) is chilled to a temperature between 0°C and 15°C outside of the deposition station and the chilled substrate is passed through the deposition station using a coating drum cooled to a temperature between 0°C and 15°C wherein the acrylate monomer is a fluorinated acrylate. 55
8. A method according to Claim 7, wherein the substrate comprises a flexible sheet and comprising the

steps of:

engaging a face of the flexible sheet with a chilled roller (42); and depositing the acrylate on the same face that engaged the chilled roller (42).

9. A method according to either one of the Claims 7 to 8, wherein the substrate comprises a plastic film and comprising the steps of:

chilling a roll of plastic film;

unrolling the plastic film;

engaging the plastic film with a chilled drum; and

rotating the drum past the deposition station.

10. A method according to either one of Claims 7 or 8, wherein the substrate comprises a flexible sheet film and comprising the steps of:

engaging a first face of the sheet with a chilled roller;

thereafter engaging the other face of the sheet with a chilled drum; and

rotating the drum past the deposition station for depositing acrylate on the first face.

11. A method according to any one of Claims 7 to 10, wherein the acrylate monomer is a polyfunctional acrylate prepolymer and the evaporating step comprises the steps of:

preheating the polyfunctional acrylate prepolymer to a temperature above ambient temperature and lower than a temperature where the prepolymer polymerizes;

atomizing the preheated prepolymer into a heated vaporization chamber in a vacuum system; and

vaporizing the prepolymer in the vaporization chamber and discharging the vaporized prepolymer into the vacuum system.

Patentansprüche

1. Verfahren zum Herstellen eines mit einer Acrylatschicht beschichteten Substrats (123), umfassend die Schritte des Verdampfens eines Acrylatmonomers, welches ein durchschnittliches Molekulargewicht im Bereich von 200 bis 600 aufweist, an einer Bedampfungsstation (131), Kondensierenlassen des Acrylatmonomers auf dem Substrat (123) als Acrylatfilm, Polymerisieren des Acrylatfilms zur Ausbildung einer polymerisierten Acrylatschicht, dadurch gekennzeichnet, daß das Substrat (123) außerhalb der Bedampfungsstation (131) auf eine Temperatur zwischen 0°C und 15°C abgekühlt wird und das abgekühlte Substrat unter Verwendung einer Beschichtungstrommel, die auf eine Temperatur zwischen 0°C und 15°C gekühlt wird, durch die Bedampfungsstation (131) geleitet wird.

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wicht im Bereich von 200 bis 600 aufweist, an einer Bedampfungsstation (131), Kondensierenlassen des Acrylatmonomers auf dem Substrat (123) als Acrylatfilm, Polymerisieren des Acrylatfilms zur Ausbildung einer polymerisierten Acrylatschicht, dadurch gekennzeichnet, daß das Substrat (123) außerhalb der Bedampfungsstation (131) auf eine Temperatur zwischen 0°C und 15°C abgekühlt wird und das abgekühlte Substrat unter Verwendung einer Beschichtungstrommel, die auf eine Temperatur zwischen 0°C und 15°C gekühlt wird, durch die Bedampfungsstation (131) geleitet wird.

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2. Verfahren nach Anspruch 1, in welchem das Substrat eine flexible Lage aufweist und umfassend die Schritte des:

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in Kontaktbringens einer Seite der flexiblen Lage mit einer abgekühlten Rolle (42); und Abscheidens des Acrylats auf derselben Seite, die die abgekühlte Rolle (42) kontaktierte.

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3. Verfahren nach einem der Ansprüche 1 bis 2, in welchem das Substrat eine Kunststoff-Folie aufweist und umfassend die Schritte des:

25

Abkühlens einer Rolle einer Kunststoff-Folie, Abrollens der Kunststoff-Folie, in Kontaktbringens der Kunststoff-Folie mit einer abgekühlten Trommel und des Rotierens der Trommel an der Bedampfungsstation (131) vorbei.

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4. Verfahren nach einem der Ansprüche 1 oder 2, in welchem das Substrat eine flexible Folienlage aufweist und umfassend die Schritte des:

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in Kontaktbringens einer ersten Seite der Lage mit einer abgekühlten Rolle (42); des darauffolgenden in Kontaktbringens der anderen Seite der Lage mit einer abgekühlten Trommel; und Abscheidung des Acrylats auf der ersten Seite.

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5. Verfahren nach einem der Ansprüche 1 bis 4, in welchem das Acrylatmonomer ein polyfunktionelles Acryl-Polymer ist und der Verdampfungsschritt die Schritte umfaßt:

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Vorheizen des polyfunktionellen Acryl-Polymer auf eine Temperatur oberhalb der Umgebungstemperatur und niedriger als eine Temperatur, bei der das Polymer polymerisiert; Zerstäuben des vorgeheizten Polymeren in eine geheizte Verdampfungskammer in einem Vakuumsystem; und Verdampfen des Polymeren in der Verdampfungskammer und Austragen des verdampften

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Prepolymers in das Vakuumsystem.

6. Verfahren nach einem der Ansprüche 1 bis 4, in welchem das Acrylat eine Mischung aus Acrylatmonomeren ist, die zumindest ein Monomer umfassen, welches ein Molekulargewicht im Bereich von 200 bis 400 aufweist, und ein Acrylat, welches ein Molekulargewicht größer als 600 aufweist. 5

7. Verfahren zum Herstellen eines mit einer Acrylatschicht beschichteten Substrats (123), umfassend die Schritte des Verdampfens eines Acrylatmonomers, welches ein durchschnittliches Molekulargewicht im Bereich von 400 bis 3000 aufweist, an einer Bedampfungsstation (131), Kondensierenlassen des Acrylatmonomers auf dem Substrat (123) als Acrylatfilm, Polymerisieren des Acrylatfilms zur Ausbildung einer polymerisierten Acrylatschicht, **dadurch gekennzeichnet, daß das Substrat (123) außerhalb der Bedampfungsstation (131) auf eine Temperatur zwischen 0°C und 15 °C abgekühlt wird und das abgekühlte Substrat unter Verwendung einer Beschichtungstrommel, die auf eine Temperatur zwischen 0°C und 15 °C gekühlt wird, durch die Bedampfungsstation (131) geleitet wird, wobei das Acrylatmonomer ein fluoriertes Acrylat ist.** 10

8. Verfahren nach Anspruch 7, in welchem das Substrat eine flexible Lage aufweist und umfassend die Schritte des: 15

(42), und Abscheidens des Acrylats auf derselben Seite, die die abgekühlte Rolle (42) kontaktierte. 20

9. Verfahren nach einem der Ansprüche 7 bis 8, in welchem das Substrat eine Kunststoff-Folie aufweist und umfassend die Schritte des: 25

Abkühlens einer Rolle einer Kunststoff-Folie, Abrollens der Kunststoff-Folie, in Kontaktbringens der Kunststoff-Folie mit einer abgekühlten Trommel; und des Rotierens der Trommel an der Verdampfungsstation vorbei. 30

10. Verfahren nach einem der Ansprüche 7 oder 8, in welchem das Substrat eine flexible Folienlage aufweist und umfassend die Schritte des: 35

in Kontaktbringens einer ersten Seite der Lage mit einer abgekühlten Rolle (42); des darauffolgenden in Kontaktbringens der anderen Seite der Lage mit einer abgekühlten Trommel; und des Rotierens der Trommel an der Bedampfungsstation vorbei zur Abscheidung von Acrylat auf der ersten Seite. 40

11. Verfahren nach einem der Ansprüche 7 bis 10, in welchem das Acrylatmonomer ein polyfunktionelles Acrylat-Prepolymer ist und der Verdampfungs- schritt die Schritte umfaßt: 45

Vorheizen des polyfunktionellen Acrylat-Prepolymers auf eine Temperatur oberhalb der Umgebungstemperatur und niedriger als eine Temperatur, bei der das Prepolymer polymerisiert; Zerstäuben des vorgeheizten Prepolymers in eine geheizte Verdampfungskammer in einem Vakuumsystem; und Verdampfen des Prepolymers in der Verdampfungskammer und Austragen des verdampften Prepolymers in das Vakuumsystem. 50

Revendications

1. Procédé de formation d'un substrat (123) revêtu d'une couche d'acrylate, comprenant les étapes consistant à évaporer un monomère d'acrylate, présentant un poids moléculaire moyen dans la plage de 200 à 600, à un poste de dépôt (131), condenser le monomère d'acrylate sur le substrat (123) sous forme d'un film d'acrylate, polymériser le film d'acrylate en vue de former une couche d'acrylate polymérisé, **caractérisé en ce que** le substrat (123) est refroidi à une température comprise entre 0 °C et 15 °C à l'extérieur du poste de dépôt (131), et en ce que le substrat refroidi est passé au travers du poste de dépôt (131), en utilisant un tambour de revêtement refroidi à une température comprise entre 0 °C et 15 °C. 55

2. Procédé selon la revendication 1, dans lequel le substrat est constitué d'une feuille souple, et comprenant les étapes consistant à :

mettre en contact une face de la feuille souple avec un rouleau refroidi (42), et déposer l'acrylate sur la face même qui a été en contact avec le rouleau refroidi (42). 60

3. Procédé selon l'une des revendications 1 ou 2, dans lequel le substrat est constitué d'un film de matière plastique, et comprenant les étapes consistant à :

refroidir un rouleau de film de matière plastique, dérouler le film de matière plastique, mettre en contact le film de matière plastique avec un tambour refroidi, et faire tourner le tambour devant le poste de dépôt (131). 65

4. Procédé selon l'une des revendications 1 ou 2, dans lequel le substrat est constitué d'un film de feuille

souple, et comprenant les étapes consistant à :

mettre en contact une première face de la feuille avec un rouleau refroidi (42),
mettre ensuite en contact l'autre face de la feuille avec un tambour refroidi, et faire tourner le tambour devant le poste de dépôt,
en vue de déposer l'acrylate sur la première face.

5. Procédé selon l'une quelconque des revendications 1 à 4, dans lequel le monomère d'acrylate est un prépolymère d'acrylate plurifonctionnel, et l'étape d'évaporation comprend les étapes consistant à :

préchauffer le prépolymère d'acrylate plurifonctionnel à une température supérieure à la température ambiante et inférieure à une température où le prépolymère se polymérise, atomiser le prépolymère préchauffé dans une chambre de vaporisation chauffée dans un système à vide, et vaporiser le prépolymère dans la chambre de vaporisation et refouler le prépolymère vaporisé dans le système à vide.

6. Procédé selon l'une quelconque des revendications 1 à 4, dans lequel l'acrylate est un mélange de monomères d'acrylates comprenant au moins un monomère_présentant_un_poids_moléculaire_dans_la_plage_de_200_à_400, et un acrylate présentant un poids moléculaire de plus de 600.

7. Procédé de formation d'un substrat (123) revêtu d'une couche d'acrylate, comprenant les étapes consistant à évaporer un monomère d'acrylate, présentant un poids moléculaire moyen dans la plage de 400 à 3 000, à un poste de dépôt (131), condenser le monomère d'acrylate sur le substrat (123) sous forme d'un film d'acrylate, polymériser le film d'acrylate en vue de former une couche d'acrylate polymérisé, **caractérisé en ce que** le substrat (123) est refroidi à une température comprise entre 0 °C et 15 °C à l'extérieur du poste de dépôt, et **en ce que** le substrat refroidi est passé au travers du poste de dépôt en utilisant un tambour de revêtement refroidi à une température comprise entre 0 °C et 15 °C, dans lequel le monomère d'acrylate est un acrylate fluoré.

8. Procédé selon la revendication 7, dans lequel le substrat est constitué d'une feuille souple, et comprenant les étapes consistant à :

mettre en contact une face de la feuille souple avec un rouleau refroidi (42), et déposer l'acrylate sur la face même qui a été en contact

avec le rouleau refroidi (42).

9. Procédé selon l'une des revendications 7 ou 8, dans lequel le substrat est constitué d'un film de matière plastique, et comprenant les étapes consistant à :

refroidir un rouleau de film de matière plastique,
déraouler le film de matière plastique,
mettre en contact le film de matière plastique avec un tambour refroidi, et faire tourner le tambour devant le poste de dépôt.

10. Procédé selon l'une des revendications 7 ou 8, dans lequel le substrat est constitué d'un film de feuille souple, et comprenant les étapes consistant à :

mettre en contact une première face de la feuille avec un rouleau refroidi,
mettre ensuite en contact l'autre face de la feuille avec un tambour refroidi, et faire tourner le tambour devant le poste de dépôt,
en vue de déposer l'acrylate sur la première face.

11. Procédé selon l'une quelconque des revendications 7 à 10, dans lequel le monomère d'acrylate est un prépolymère d'acrylate plurifonctionnel, et l'étape d'évaporation comprend les étapes consistant à :

préchauffer le prépolymère d'acrylate plurifonctionnel à une température supérieure à la température ambiante et inférieure à une température où le prépolymère se polymérise, atomiser le prépolymère préchauffé dans une chambre de vaporisation chauffée dans un système à vide, et vaporiser le prépolymère dans la chambre de vaporisation et refouler le prépolymère vaporisé dans le système à vide.

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Fig. 1

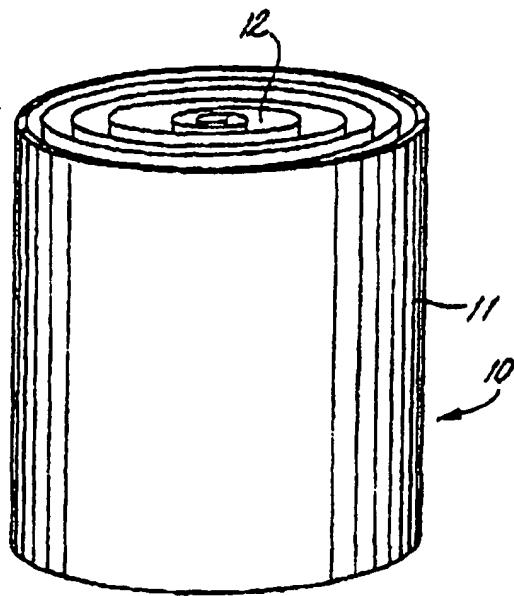


Fig. 2a

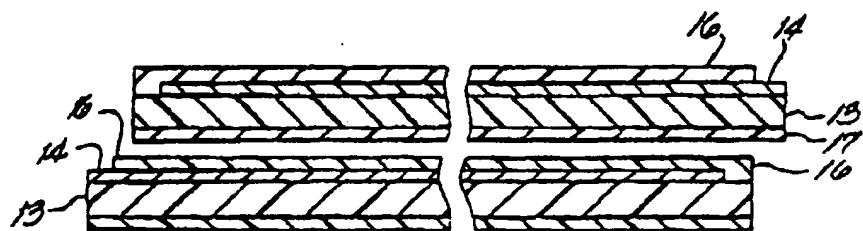


Fig. 2b



Fig. 2c



Fig. 3

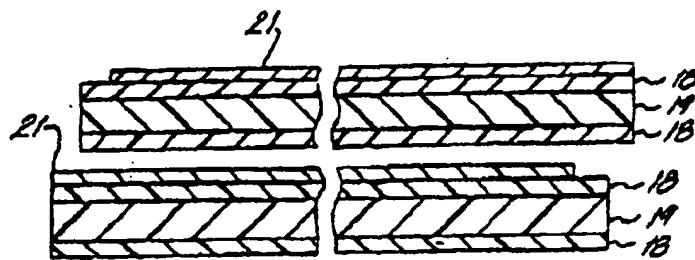


Fig. 4

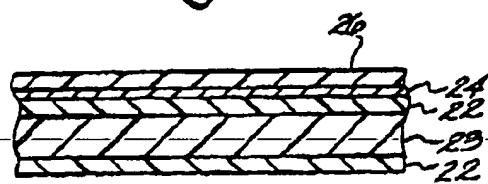
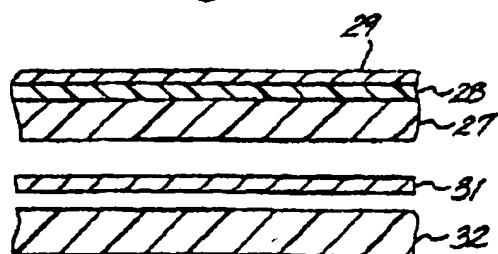


Fig. 5



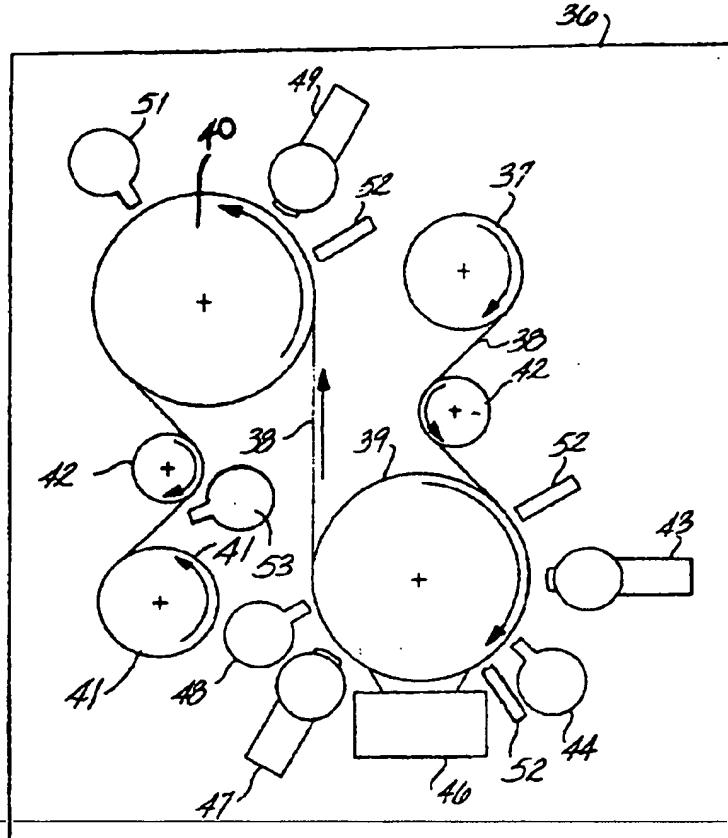


Fig. 6

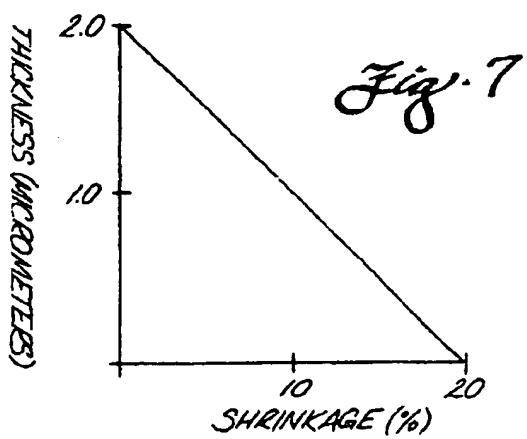


Fig. 8

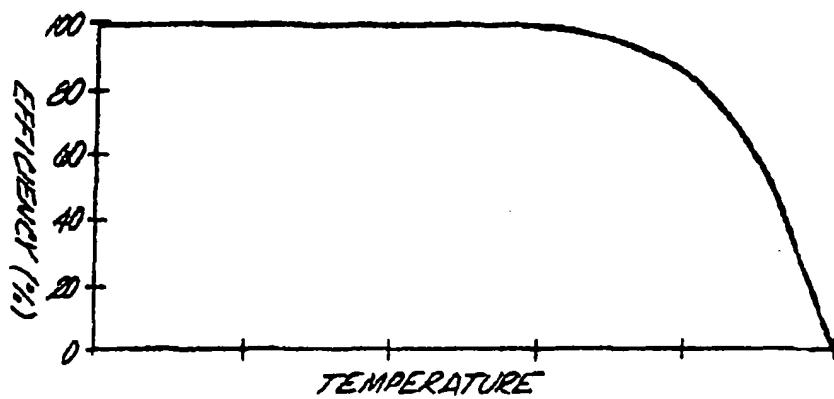


Fig. 9

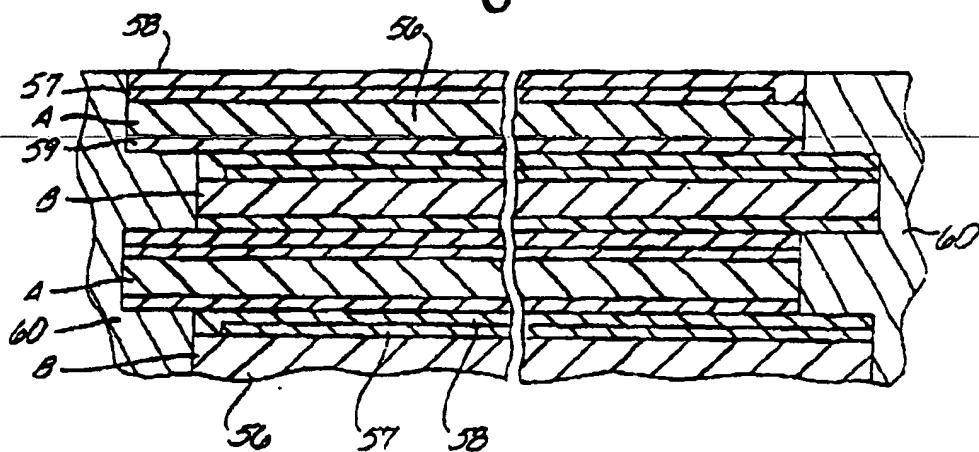


Fig. 10

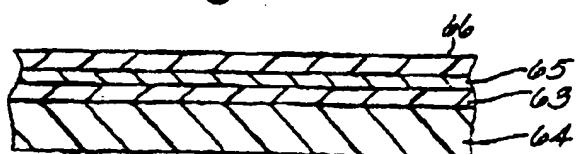


Fig. 11

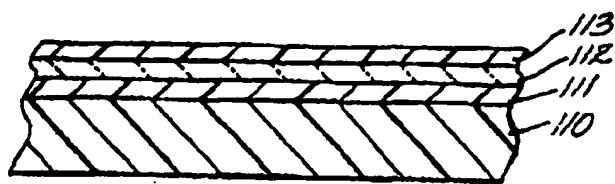


Fig. 12

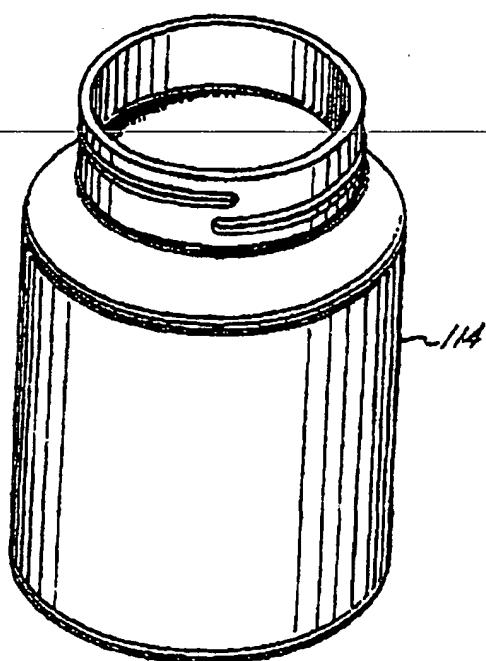


Fig. 13

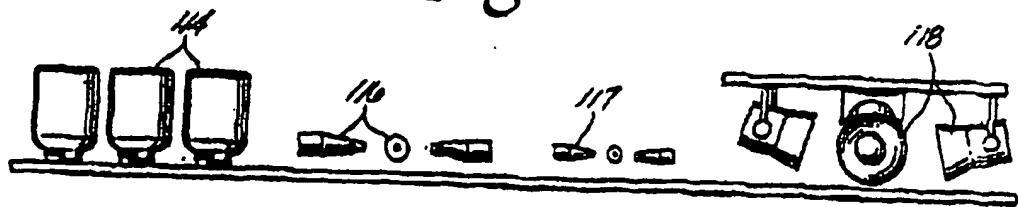


Fig. 14

